

Prisca 5.1.0.17
Date of report: 14-06-2025

NA

Patient data			
Name	Mrs. AARTI CHAVHAN		Patient ID 0372506130091
Birthday	29-12-1996		Sample ID B2917297
Age at sample date	28.5		Sample Date 13-06-2025
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	79	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 12 + 5
PAPP-A	1.49 mIU/mL	0.44	Method CRL Robinson
fb-hCG	36.58 ng/mL	1.07	Scan date 11-06-2025
Risks at sampling date			Crown rump length in mm 66
Age risk	1:776		Nuchal translucency MoM 0.89
Biochemical T21 risk	1:544		Nasal bone present
Combined trisomy 21 risk	1:3083		Sonographer NA
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT NA
Risk			Trisomy 21
			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 3083 women with the same data, there is one woman with a trisomy 21 pregnancy and 3082 women with not affected pregnancies. The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that risk calculations are statistical approaches and have no diagnostic value! The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)). The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!
Trisomy 13/18 + NT			
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			

Sign of Physician